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09/926596

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(21) Patentansökningsnummer 9901961-4  
Patent application number

(86) Ingivningsdatum 1999-05-28  
Date of filing

Stockholm, 2000-07-24

För Patent- och registreringsverket  
For the Patent- and Registration Office

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### Field of the invention

The present invention is based on the finding that two fimbrial operons, the *saf* operon and the *tcf* operon, are specific for *Salmonella enterica* subspecies I bacteria and therefore have therapeutic use. Due to their specificity they can be used to provide vaccines against *Salmonella enterica* subspecies I as well as for detection of *Salmonella enterica* subspecies I. The *saf* operon is specific for all *Salmonella enterica* subspecies I bacteria and the *tcf* operon is specific for the serovar Typhi of *Salmonella enterica* subspecies I.

- 10 All or part of the DNA-sequences of the genes encoding these proteins can be used as active agents in a vaccine against diseases caused by the *Salmonella enterica* subspecies I bacterial strains or for detection of said bacterial strains.

The present invention also relates to methods of isolating these fimbrial proteins, to antibodies directed against these proteins, and to a vaccine composition comprising these proteins or antibodies directed against these proteins for use in the treatment of infections caused by the *Salmonella spp.* The fimbrial proteins according to the invention or antibodies directed against them can be used for detection of *Salmonella spp.* bacteria.

20

### Background of the invention

The members of genus *Salmonella spp.* colonize and infect a wide range of different organisms. Many cause gastroenteritis and enteric fever in humans and domesticated animals while others are not associated with human disease (Saylers et al, 1994). The genus has been divided into two species, *Salmonella bongori* and *Salmonella enterica* where *enterica* can be further subdivided into seven subspecies, designated I, II, IIIa, IIIb, IV, VI, and VII (Reeves et al, 1989). *Salmonella enterica* subspecies I are preferentially associated with warm-blooded animals. Over 99% of all clinical *Salmonella* isolates are strains belonging to this subspecies, including serovars Typhimurium and Enteritidis, which are the major causes of *Salmonella* induced gastroenteritis in humans, and Typhi, the human specific causative organism of typhoid fever, the most severe form of human salmonellosis (Popoff et al, 1992).

35 *Salmonella enterica* subspecies I consists of over 1300 different serovars and is preferentially associated with warm-blooded animals (Baumler, 1994). Over 99% of all clinical *Salmonella* isolate are strains belonging to this subspecies,

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including serovars Typhimurium and Enteritidis, which are the major causes of *Salmonella* induced gastroenteritis in humans, and Typhi, the human specific causative organism of typhoid fever, the most severe form of human salmonellosis (Popoff and Le Minor, 1992).

5

Today gastroenteritis and enteric fever can neither be prevented nor treated with good results. Typhoid fever is a substantial public health problem in developing countries. Each year 33 million people become ill and over 500 000 people die from this infection (American Institute of Medicine, 1986). Typhoid fever can be prevented by vaccination with attenuated bacteria, such as Ty21 and Vi vaccines and whole cell vaccines. Whole cell vaccines show a high incidence of side effects (Ashcroft et al, 1964, Yugoslav Typhoid commission, 1964). The vaccines consisting of attenuated strains of *Salmonella typhi* suffer from serious drawbacks. They must be administered as three or four spaced doses in order to stimulate protective immune responses (Levine et al, 1989). The treatment of *Salmonella typhi* with antibiotics is jeopardized since there are strains of *Salmonella typhi* that are resistant to chloramphenicol, ampicillin, and trimethoprim as well as ciprofloxacin (i.e. multidrug-resistant strains) (Rowe et al, 1997).

20

Accurate detection of *Salmonella enterica* subspecies I is today not possible. *Salmonella enterica* subspecies I can today only be detected by antibodies directed against surface proteins of *Salmonella enterica* subspecies I. The use of the sequences according to the invention makes it for the first time possible to rapidly and accurately determine the presence of *Salmonella enterica* subspecies I.

25

For many pathogenic bacteria, there is evidence that the filamentous surface protein structures called pili (fimbriae) are connected to the adhesion of the bacteria to the host cells. Pili proteins are very antigenic and are easily purified. Therefore pili preparations have been used as antigens for vaccination.

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#### Summary of the invention

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The invention relates to the objects as defined in the claims. The main object of the present invention is to provide two fimbrial proteins that are specific for *Salmonella enterica* subspecies I bacterial strains, the nucleotide sequences

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## 30

**Figure 1.**

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The clones were selected from partial *Eco* RI and *Bam*HI libraries in the Lambda Dash II vector. The cs7 insert is represented by a bold line. The extent

of respective phage insert is represented by horizontal bars. Name and size of the phage inserts are indicated on the left side of the figure.

Figure 2.

Schematic representation of the pTY52 cosmid comprising the *tcf*-operon (SEQ ID NO 2).

- 5 A *tcf* specific PCR fragment of 11105 bp was cloned into the Expand vector I cosmid (Roche). The insert is represented with a thick black line while vector sequences are represented with thin lines. Relevant restriction sites sequences are indicated. The position of the *tcf*-operon, i.e. *tcfA*, *B*, *C* and *D* (SEQ ID NO 2), is represented by a shaded arrow.

10 Figure 3.

The phylogenetic distribution of the identified genes on the cs7 insert was investigated using the well defined SARC collection, see Example 1.

Figure 4.

- 15 A 2 kb large internal *EcoR* I fragment was used as a probe in a Southern blot of the SARC collection, see Example 2.

#### Sequence listing

- 20 SEQ ID NO 1—DNA sequence of the genes encoding the precursor of the *saf* fimbrie unit of *Salmonella enterica* subspecies I.

SEQ ID NO 2—DNA sequence of the genes which encode the precursor of the *tcf* fimbrie unit of *Salmonella enterica* subspecies I serovar Typhi.

#### Deposit information

- 25 The phages carrying the inserted SEQ ID NO 1, i.e. phages clones B1, D1, F11 and N10 (see Figure 1) have been given the ECACC Accession numbers 99051922, 99051923, 99051924, and 99051925, respectively.

The cosmide carrying the inserted SEQ ID NO 2, i.e. cosmide pTY52 (see Figure 2) has been given the ECACC Accession number 99051926.

- 30 The depositions were made May 19, 1999.

#### Detailed description of the invention

- The present invention is based on the finding that two fimbrial operons, the *saf* operon and the *tcf* operon, are specific for *Salmonella enterica* subspecies I bacteria. Due to their specificity they can be used to provide vaccines against *Salmonella enterica* subspecies I as well as detection methods for *Salmonella enterica* subspecies I. The *saf* operon is specific for all *Salmonella enterica*
- 35

subspecies 1 bacteria and the *tcf* operon is specific for the serovar typhi of *Salmonella enterica* subspecies 1, see Examples 1 & 2.

5 The main object of the invention relates to two fimbrial operons, the *saf* operon and the *tcf* operon, that are specific for *Salmonella enterica* subspecies 1 bacteria for therapeutic use.

Another object of the present invention is to provide vaccines against *Salmonella enterica* subspecies 1 induced gastroenteritis, enteric fever and  
10 typhoid fever.

A further object of the present invention is to provide methods to detect *Salmonella enterica* subspecies 1. The nucleotide sequences according to the invention are useful for constructing vectors for use as vaccines for insertion  
15 into attenuated bacteria in constructing a recombinant vaccine, for insertion into a viral vector in constructing a recombinant viral vaccine, or for direct inoculation as a nucleic acid vaccine. The pili proteins according to the invention, or antigenic fragments thereof, can be used for active immunization and antibodies directed against them can be used for passive immunization. All  
20 these applications of the sequences according to the invention are obtained by applying standard techniques known to the man ordinary skilled in the art.

Vaccines against *Salmonella enterica* subspecies 1.

The genes encoding the *saf* and *tcf* fimbrial structures, or fragments thereof,  
25 may be incorporated into a bacterial or viral vaccine comprising recombinant bacteria, virus or fungi which are engineered to produce one or more immunogenic epitopes of the *saf* or *tcf* fimbrial structures. In addition, the genes encoding the *saf* and *tcf* fimbrial structures, or part thereof, operatively linked to regulatory elements, can be introduced directly as a nucleic acid  
30 vaccine, to elicit a protective immune response.

The proteins or antigenic fragment thereof, deduced from the nucleic acid sequences of the present invention are useful alone or in conventional vaccine mixtures in the vaccine compositions according to the invention. The proteins  
35 could be produced by chemical synthesis or recombinant expression according to conventional methods.

The proteins and peptides according to the invention can be obtained by using a host organism transformed or transfected with an expression vector obtained by insertion of a gene according to the invention, or part thereof, into a vector in a conventional manner. The vector which is used to construct the expression  
5 vector is not particularly limited, but specific examples include plasmids such as pET (Stratagen) and the like; and phages such as M13 (NEB), phage display libraries and the like. As expression regulatory sequence can among others T7 promoters and lac promoters be used.

10 An appropriate host to be transformed or transfected with the expression vector can be chosen among for example *E.-coli*, *Salmonella* or *Bacillus subtilis*. The transformed or transfected host is cultured and proliferated under suitable conditions.

15 After culturing, the peptides of the present invention may be purified by, for example, chromatography, precipitation, and/or density gradient centrifugation. The thus obtained peptides can be used as a vaccine or for the production of antibodies directed against said peptides, which can be used for passive immunization.

20 The purified preparation containing one or several proteins according to the invention, or parts thereof, is then formulated as a pharmaceutical composition, as for example a vaccine, or in a mixture with adjuvants. If desired the proteins are fragmented by standard chemical or enzymatic  
25 techniques to produce antigenic segments.

In formulating the vaccine compositions with the peptide or protein, alone or in various combinations, the immunogen is adjusted to an appropriate concentration and formulated with any suitable vaccine adjuvant. The  
30 immunogen may also be incorporated into liposomes, or conjugated to polysaccharides and/or other polymers for use in a vaccine formulation.

The different vaccines according to the present invention are administered to mammals in many different ways. These include intradermal, intramuscular,  
35 intraperitoneal, intravenous, subcutaneous, oral, and intranasal routes of administration. The vaccine doses will differ depending on circumstances such

as body weight, interferences with other administered medicaments etc. The upper limit is not critical unless the dose shows toxicity.

The peptides and proteins of the present invention are also useful to produce  
 5 monoclonal or polyclonal antibodies for use in passive immunotherapy against *Salmonella enterica* subspecies 1. Human immunoglobulin is preferred. Antisera is obtained from individuals immunized with proteins or peptides according to the invention. The immunoglobulin fraction is then enriched, for example by immunoaffinity or affinity chromatography. Antibodies raised in  
 10 a suitable mammal or in the patient to be treated, can subsequently be administered locally or topically, e.g. orally to the patient.

Detection of *Salmonella enterica* subspecies I in general.

The sequences according to the invention, or part thereof, or fragments  
 15 hybridizing therewith, as well as the proteins according to the invention, or part thereof, and antibodies directed to said proteins, or antigenic fragments thereof, can be used in molecular diagnostic assays for the detection of *Salmonell enterica* subspecies I.

20 Nucleic acids having the nucleotide sequence according to the invention, or any nucleotide sequence hybridizing therewith can be used as a probe in nucleic acid hybridization assays for the detection of *Salmonella spp* in various tissues and body fluids of patients. The hybridization assay may be of any type including; Southern blots, Northern blots, colony blots.

25 PCR technology is the most preferred technology for detection according to the invention of *Salmonella enterica* subspecies 1. Primers of at least one selected from the 5' end and one from the 3' end can be used in PCR and other known tests to rapidly identify the presence of *Salmonella enterica* subspecies 1. This is  
 30 according to conventional techniques.

The isolated and purified proteins and peptides of the invention can be used as  
 35 diagnostics to measure an increase in serum titer of *Salmonella enterica* subspecies I-specific antibody since they bind strongly to these antibodies. A serum test sample can be screened for *Salmonella enterica* subspecies I by methods such as for example ELISA.



The invention further comprises the use of antibodies directed against the *saf* and *tcf* fimbriae structures for quantitative or qualitative determinations of the pili proteins of the invention, or fractions thereof, in cells, tissues or body fluids.

5

Detection of *Salmonella enterica* subspecies I by using nucleic acid hybridization technology

Nucleic acid hybridization technology can also be to detect *Salmonella enterica* subspecies I according to the invention. The nucleic acid probes chosen from  
10 parts of the sequences according to the invention can be either DNA or RNA. DNA sequences complementary to the sequences according to the invention can also be used. The binding of the probe to the target sequence, i.e. the hybridization, must not be perfect. Variations and mutations of the sequences according to the invention can be used as long as they hybridize good enough  
15 to detect *Salmonella enterica* subspecies I. The preferred length of the nucleic acid probes is about 10 to 400 nucleotides, most preferred not longer than 100 nucleotides.

The nucleotide probe is preferably chosen from the parts of the sequences that  
20 have the least variation. In the most preferred embodiments when screening for SEQ ID NO 1 (the *saf* operon, specific for *Salmonella enterica* subspecies I) a nucleotide probe or PCR primer selected from nucleotides 37 368-37 868 should be avoided since this region is hypervariable.

25 The nucleic acid probes according to the invention are prepared by any conventional method such as organic synthesis, recombinant techniques, or isolation from genomic DNA.

30 The nucleic acid probes of the invention are labeled in a conventional manner to signal hybridization to target nucleic acid from *Salmonella enterica* subspecies I. The labeling may comprise a radiolabel, an enzyme, a bacterial label, a fluorescent label, an antibody, an antigen, a latex particle, an electron dense compound, or a light scattering particle.

35 The probes may be provided in a lyophilized form, to be reconstituted in a buffer appropriate for hybridization, or the probes may already be present in

such a buffer. The buffer may contain a suitable hybridization enhancer, detergent, carrier DNA, and a compound to increase the specificity.

Any conventional hybridization assay technique, such as dot blot hybridization, Southern blotting, sandwich hybridization, displacement hybridization and the like, can be used.

The target analyte polynucleotide of a microorganism may be in various media, most often in a biological, or physiological specimen. In most cases it is preferred to subject the specimen containing the target polynucleotide to any conventional extraction, purification, and/or isolation before conducting the analysis.

The sample containing the target analyte nucleotide sequence must often be treated to convert the DNA to a single-stranded form, which may be accomplished by a variety of conventional techniques, such as thermal or chemical techniques.

The following examples describe the isolation and specificity of the sequences according to the invention.

#### EXAMPLE 1

Identification and characterization of the *saf* operon.

The present inventors found, upon investigation of a 7 kb chromosomal region on centisome 7 originally isolated from the *S. typhimurium* strain SR-11 k3181, a region that exhibits many of the traits that define a pathogenicity island. It has a lower G+C composition than the average composition of the *Salmonella* genome and includes many sequences related to different mobile genetic elements. The region is not present in *E. coli* K12, and the *Salmonella* specific DNA is inserted between the tRNA gene *aspV* and the stop codon of *yafV*, a hypothetical protein upstream of the *yafH* gene at 5 min in the *E. coli* chromosome. This *Salmonella* specific insert encodes proteins creating adhesive structures and other virulence factors. Sequencing revealed genes encoding a new fimbrial operon that they designated *Salmonella* Atypical Fimbriae (*saf*), due to its relatedness to a subgroup of adhesive structures forming thin atypical fimbriae or non-fimbrial adhesins.

The *saf* operon consists of four contiguous genes, *safA*, *safB*, *safC* and *safD* that encode fimbrial subunit, periplasmic chaperone, outer membrane usher protein and alternative fimbrial subunit, respectively. The genes *safA*, *B*, *C* and *safD* encode putative proteins of 166, 244, 836 and 156 amino acids, respectively. Analyzes of clinical *Salmonella* isolates showed that DNA of 195 out of 198 clinical isolates belonging to *S. enterica* subspecies I hybridized with *safB* and *safC*, i.e. these sequences are common to more than 99% of the known *Salmonella enterica* subspecies 1 bacteria. The inventors showed that 58% of these clinical isolates carry the *safA*, see Table 1.

Table 1. The prevalence of the *saf* genes in clinical *Salmonella* isolates.

Serovar	<i>safA</i>	<i>safB</i>	<i>safC</i>	# isolates
<i>S. adelaide</i>	-	+	+	1
<i>S. agona</i>	+	+	+	6
<i>S. anatum</i>	-	+	+	3
<i>S. bareilly</i>	+	+	+	3
<i>S. blockley</i>	+	+	+	3
<i>S. bovismorbificans</i>	-	+	+	5
<i>S. braenderup</i>	-	+	+	4
<i>S. brandenburg</i>	+	+	+	1
<i>S. bredeney</i>	+/-	+	+	15
<i>S. chester</i>	+	+	+	1
<i>S. colindale</i>	-	+	+	1
<i>S. derby</i>	-	+	+	1
<i>S. dublin</i>	-	+	+	1
<i>S. eastbourne</i>	+	+	+	2
<i>S. emek</i>	+	+	+	1
<i>S. enteritidis</i>	-	+	+	8
<i>S. give</i>	-	+	+	1
<i>S. goettingen</i>	+	+	+	1
<i>S. haardt</i>	-	+	+	1
<i>S. hadar</i>	+	+	+	16
<i>S. heidelberg</i>	-	+	+	1
<i>S. huttingfoss</i>	+	+	+	5
<i>S. infantis</i>	-/+	+	+	6
<i>S. java</i>	-	+	+	1
<i>S. javiana</i>	-	+	+	1
<i>S. kottbus</i>	-	+	+	1
<i>S. livingstone</i>	-	+	+	1
<i>S. london</i>	+	+	+	1
<i>S. maastricht</i>	+	+	+	2
<i>S. mbandaka</i>	-	-	-	3
<i>S. montevideo</i>	+	+	+	1
<i>S. muenster</i>	-	+	+	1
<i>S. newport</i>	+	+	+	2
<i>S. ohio</i>	+	+	+	1
<i>S. oranienburg</i>	+	+	+	2
<i>S. panama</i>	+	+	+	3
<i>S. potsdam</i>	+	+	+	1
<i>S. rissen</i>	-	-	-	1
<i>S. saarbrücken</i>	-	+	+	1
<i>S. saint paul</i>	+	+	+	3
<i>S. schwarzengrund</i>	-	+	+	1
<i>S. singapore</i>	+	+	+	1
<i>S. stanley</i>	+	+	+	5
<i>S. subsp I 4.5, 12:-:-</i>	+	+	+	2
<i>S. subsp I 4.5, 12:b:-</i>	-	+	+	1
<i>S. subsp I 4.5, 12:i:-</i>	+	+	+	1
<i>S. subsp I spont</i>	-	+	+	1
<i>S. tennessee</i>	+	+	+	2
<i>S. thompson</i>	-	+	+	1
<i>S. typhi</i>	-	+	+	1
<i>S. typhimurium</i>	+	+	+	27
<i>S. virchow</i>	+	+	+	7
<i>S. weltervreden</i>	-	+	+	1
<i>S. worthington</i>	-	-	-	2
<i>S. subsp III</i>	-	-	-	1

The phylogenetic distribution of the identified genes on the cs7 insert was investigated using the well defined SARC collection, which showed that the presence of the *safA*, *safB*, *safC* and *safD* genes is restricted to *S. enterica* subspecies I (Fig. 3). This region is hence the first subspecies I specific genetic region to be identified with a broad distribution within the subspecies. Since the serovars of subspecies I constitute over 99% of human salmonellosis and are preferentially associated with warm blooded animals, it implicates a role for the *saf* adhesive organelle in the colonization of these organisms.

## EXAMPLE 2

Identification and characterization of the *tcf* operon.

The present inventors found that *Salmonella enterica* subspecies I serovar Typhi contains DNA encoding an additional fimbrial operon, the *tcf* operon, in the *sinR-pagN* intergenic region. Southern blot analysis revealed a markedly different restriction pattern in *S. enterica* serovar Typhi than the other subspecies I isolates, suggesting that the *saf-sin* region in serovar Typhi might carry additional DNA relative to serovar Typhimurium strains. A PCR reaction (using a kit from Roche) was therefore performed using a *sinR* (5'-GTA AAT CGC TTA GTC GCC-3') specific forward primer and a *pagN* (5'-TCA ACT CAA CCT TCA GCC-3') specific reverse primer.

This primer pair produced, as expected, a product of 2 kb in serovar Typhimurium from the SARC collection, while from serovar Typhi the product was 10 kb. Thus, the neighboring *sinR* and *pagN* genes in serovar Typhimurium strains are separated by approximately 8 kb in serovar Typhi.

The Typhi specific PCR product was purified, digested partially with *EcoRI* and sub-cloned into pUC18 forming a set of overlapping clones. Sequencing of the clones revealed a putative fimbrial operon designated *tcf* for Typhi Colonizing Factor. Four ORFs, *tcfA,B,C,D*, have been identified with putative proteins having significant homology to CooB (38% identical over 192 aa), CooA (37% identical over 170 aa), CooC (34% identical over 872 aa) and CooD (31% identical over 272 aa), respectively. The Coo proteins are involved in the biosynthesis of the CS1 colonizing factor antigens of enterotoxigenic *E.coli* (Fig. 4) (Froehlich et al., 1994). The peptide of the *tcfB* ORF is also homologous to the CblA major fimbrial subunit protein (45% identical over 154 aa) of the cable

- type II pili of the cystic fibrosis-associated *Burkholderia cepacia* (Sajjan et al., 1995). Down-stream of the *tcf*-operon two ORFs were identified with the same transcriptional orientation as the *tcf* genes. The first was designated *tinR* for Typhi insert regulator because it is homologous (33% identical over 144 aa) to
- 5 *AzlB* of *Bacillus subtilis*, a member of the Lrp/AsnC family of transcriptional regulators (Belitsky et al., 1997). *tinR* is followed by an ORF (*tioA* for Typhi insert orf) encoding a putative protein of 205 amino acids with no significant homologies to anything in the DDBJ/EMBL/GenBank databases. The above
- 10 sequence from *Salmonella enterica* serovar Typhi strain RKS 3333 and the *tcf* region of the incomplete genome sequence from serovar Typhi strain CT18 ([http:// www.sanger.ac.uk](http://www.sanger.ac.uk)) are 99% identical over the total length of the investigated region in concordance with the clonal nature of the serovar .
- 15 A 2 kb large internal *EcoR* I fragment was used as a probe in a Southern blot of the SARC collection. This blot shows that *Salmonella enterica* subspecies I serovar Typhi (SARC2) is the only strain in the collection possessing DNA hybridizing to this fragment (Fig. 4).

SEQUENCE LISTING NO. 1

&lt;110&gt; Folkesson, Anders

<120> The complete sequence of the cs7 insert in Salmonella enteric serovar Typhimurium

<130> Complete sequence of the cs7 insert

<140>

<141>

<160> 5

&lt;170&gt; PatentIn Ver. 2.1

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<211> 46870

<212> DNA

<213> Salmonella typhimurium

<220>

<221> CDS

<222> (37368) .. (37868)

<223> safA putative fimbrial subunit

<220>

<221> CDS

<222> (37952) .. (38689)

<223> safB putative periplasmic chaperone

**<220>**

&lt;221&gt; CDS

<222> (38713) .. (41223)

<223> safC putative outer membrane usher

**<220>**

&lt;221&gt; CDS

<222> (41245) .. (41715)

<223> safD putative fimbrial subunit

<400> 1

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caaatgggcc tacgcctgga tgacgaacag gatattaccg ccacttcttt cactgtcatg 180

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tgggatggtc gggctactgg gttcactgct ctcaaaccag cgacgaatgc accgcaggcg 480

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<212> PRT

<213> Salmonella typhimurium

<400> 2

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Leu Ile Pro Val Ser Gly Leu Lys Ala Gly Lys Asn Ala Pro Ser Ala  
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Lys Ile Ala Lys Leu Val Val Asn Ser Thr Thr Leu Lys Glu Phe Gly  
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Val Arg Gly Ile Ser Asn Asn Val Val Asp Ser Thr Gly Thr Ala Trp  
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Arg Val Ala Gly Lys Asn Thr Gly Lys Glu Ile Gly Val Gly Leu Ser  
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Asn Trp Met Thr Phe Asn Ser Asn Asp Thr Leu Asp Ile Val Leu Thr  
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Asn Ser Met Val Ser Val Tyr Ala Val Asn Gln Gln Leu Asn Ser Ala  
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Thr Lys Leu Phe Ser Val Lys Leu Gly Ala Thr Arg Val Ile Tyr His  
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Ala Gly Thr Ala Gly Ala Thr Leu Ser Val Ser Asn Pro Gln Asn Tyr  
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Pro Ile Leu Val Gln Ser Ser Val Lys Ala Ala Asp Lys Ser Ser Pro  
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Ala Pro Phe Leu Val Met Pro Pro Leu Phe Arg Leu Glu Ala Asn Gln  
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Gln Ser Gln Leu Arg Ile Val Arg Thr Gly Gly Asp Met Pro Thr Asp  
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Arg Glu Thr Leu Gln Trp Val Cys Ile Lys Ala Val Pro Pro Glu Asn

115

120

125

Glu Pro Ser Asp Thr Gln Ala Lys Gly Ala Thr Leu Asp Leu Asn Leu  
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 Ser Ile Asn Ala Cys Asp Lys Leu Ile Phe Arg Pro Asp Ala Val Lys  
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 Gly Thr Pro Glu Asp Val Ala Gly Asn Leu Arg Trp Val Glu Thr Gly  
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 Asn Lys Leu Lys Val Glu Asn Pro Thr Pro Phe Tyr Met Asn Leu Ala  
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 Ser Val Thr Val Gly Gly Lys Pro Ile Thr Gly Leu Glu Tyr Val Pro  
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 Pro Phe Ala Asp Lys Thr Leu Asn Met Pro Gly Ser Ala His Gly Asp  
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 His Tyr Val Leu Lys  
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&lt;210&gt; 4

&lt;211&gt; 836

&lt;212&gt; PRT

&lt;213&gt; Salmonella typhimurium

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 Gln Pro Gly Thr Tyr Arg Val Asp Val Met Val Asn Gly Lys Arg Val  
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 Asp Thr Arg Asp Val Val Phe Lys Leu Glu Lys Asp Gly Gln Gly Thr  
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 Pro Val Leu Ala Pro Cys Leu Thr Val Ser Gln Leu Ser Arg Tyr Gly  
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 Val Lys Thr Glu Asp Tyr Pro Gln Leu Trp Lys Ala Ala Lys Pro Pro  
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 Asp Glu Cys Ala Asp Leu Thr Ala Ile Pro Gln Ala Lys Ala Val Leu  
 115 120 125  
 Asp Ile Asn Asn Gln Gln Leu Gln Leu Ser Ile Pro Gln Leu Ala Leu  
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 Arg Pro Glu Phe Lys Gly Ile Ala Pro Glu Asp Leu Trp Asp Asp Gly

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Tyr Lys Met Asp Met Val Gly Arg Asp Asn Ser Ser Trp Val Gln Leu			
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Gln Pro Gly Ile Asn Ile Gly Ala Trp Arg Val Arg Asn Ala Thr Ser			
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Trp Gln Arg Ser Ser Gln Leu Ser Gly Lys Trp Gln Ala Ala Tyr Thr			
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Tyr Ala Glu Arg Gly Leu Tyr Ser Leu Lys Ser Arg Leu Thr Leu Gly			
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Gln Lys Thr Ser Gln Gly Glu Ile Phe Asp Ser Val Pro Phe Thr Gly			
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Val Met Leu Ala Ser Asp Asp Asn Met Val Pro Tyr Ser Glu Arg Gln			
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Phe Ala Pro Val Val Arg Gly Ile Ala Arg Thr Gln Ala Arg Val Glu			
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Val Lys Gln Asn Gly Tyr Thr Ile Tyr Asn Thr Thr Val Ala Pro Gly			
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Pro Phe Ala Leu Arg Asp Leu Ser Val Thr Asp Ser Ser Gly Asp Leu			
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His Val Thr Val Trp Glu Ala Asp Gly Ser Thr Gln Met Phe Val Val			
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Pro Tyr Gln Thr Pro Ala Ile Ala Leu His Gln Gly Tyr Leu Lys Tyr			
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Ser Leu Leu Ala Gly Arg Tyr Arg Ser Ser Asp Ser Ala Thr Asp Lys			
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Arg Gln Ile Ala Gln Ala Thr Leu Met Tyr Gly Leu Pro Trp Asn Leu			
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Thr Ala Tyr Gly Gly Ile Gln Ser Ala Thr His Asn Gln Ala Ala Leu			
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Leu Gly Leu Gly Gly Ser Leu Gly Arg Trp Gly Ser Leu Ser Val Asp			
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Gly Ser Asp Thr His Ser Gln Arg Gln Gly Glu Ala Val Gln Gln Gly			
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Ala Ser Trp Arg Leu Arg Tyr Ser Asn Gln Leu Thr Ala Thr Gly Thr			
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Asn Phe Phe Leu Thr Arg Trp Gln Tyr Ala Ser Gln Gly Tyr Asn Thr			
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Leu Ser Asp Val Leu Asp Ser Tyr Arg His Asn Gly Asn Arg Leu Trp			
	465	470	475
			480

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Lys Gly Glu Leu Thr Val Lys Trp Gly Ala Gln Gln Cys Arg Val Asn  
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35 40 45

Val Pro Asp Gly Met Val Leu Ala Gln Gly Trp Val Thr Tyr His Gly  
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Ser His Ser Gly Phe Arg Val Trp Ser Asp Glu Gln Lys Ala Gly Asn  
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Thr Pro Thr Val Leu Leu Leu Ser Gly Gln Gln Asp Pro Arg His His  
85 90 95

Ile Gln Val Arg Leu Glu Gly Glu Gly Trp Gln Pro Asp Thr Val Ser  
100 105 110

Gly Arg Gly Ala Ile Leu Arg Thr Ala Ala Asp Asn Ala Ser Phe Ser  
115 120 125

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## SEQUENCE LISTING NO. 2

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Val Gln Lys Asp Ile Thr Val Thr Ala Asn Ile Asp Ser Thr Leu Glu			
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Phe Met Pro Gly Lys Gly Leu Val His Lys Ser Leu Gln Thr Arg Leu			
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Leu Glu Gly Gln Thr Glu Gln Ile Glu Val Leu Leu Pro Gly His Ser			
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 Ala Gly Thr Asp Ile His Ser Ser Arg Asn Gln Lys Thr Gln Thr Ser  
 1015 1020 1025 1030

tgg aat gtg aac cat gtg aga tcc tgg cag cag gat ctg tat cgt gaa 5210  
 Trp Asn Val Asn His Val Arg Ser Trp Gln Gln Asp Leu Tyr Arg Glu  
 1035 1040 1045

ctg tcg gtg ggt ttc tcc ggt tat aac gac gac agc tgg agc ggg agt 5258  
 Leu Ser Val Gly Phe Ser Gly Tyr Asn Asp Asp Ser Trp Ser Gly Ser  
 1050 1055 1060

ctc ggc gga cgc atg agc ggc cgt atg ggt gaa ctg agc gcc act atc 5306  
 Leu Gly Gly Arg Met Ser Gly Arg Met Gly Glu Leu Ser Ala Thr Ile  
 1065 1070 1075

agt aac tcc cat caa cgt aat gcg ggc agc gcc agt tca ctc acc gct 5354  
 Ser Asn Ser His Gln Arg Asn Ala Gly Ser Ala Ser Ser Leu Thr Ala  
 1080 1085 1090

ggc tac agc tcg tct ctg gcg tta tcc cgt aat gga ctg ttc tgg gga 5402  
 Gly Tyr Ser Ser Ser Leu Ala Leu Ser Arg Asn Gly Leu Phe Trp Gly  
 1095 1100 1105 1110

ggt ggt cag gac ggt gaa ccg gcc tct ggc atg gcg gtg aac gtg gag 5450  
 Gly Gly Gln Asp Gly Glu Pro Ala Ser Gly Met Ala Val Asn Val Glu  
 1115 1120 1125

tca gag ggg gac gag ggc agt agc ggg aaa gta gtc agc gtt cgt ggc 5498  
 Ser Glu Gly Asp Glu Gly Ser Ser Gly Lys Val Val Ser Val Arg Gly  
 1130 1135 1140

agc agc cag ccg ttc agt ctc ggt ttt ggt cag cag tcg ctg ttg ctg 5546  
 Ser Ser Gln Pro Phe Ser Leu Gly Phe Gly Gln Gln Ser Leu Leu Leu  
 1145 1150 1155

atg gaa ggc tat aac gcc acg gag gtg acc att gag gat gca ggg gtt 5594  
 Met Glu Gly Tyr Asn Ala Thr Glu Val Thr Ile Glu Asp Ala Gly Val  
 1160 1165 1170

agt tca cag ggt atg gca ggc gta aaa gcg gga ggg gga agc agg tgt 5642  
 Ser Ser Gln Gly Met Ala Gly Val Lys Ala Gly Gly Gly Ser Arg Cys



1175	1180	1185	1190	
tac ttc ctg aca ccc ggg cat ctg ctg gtt cac aac atc agc gcc agt				5690
Tyr Phe Leu Thr Pro Gly His Leu Leu Val His Asn Ile Ser Ala Ser				
1195		1200	1205	
atg agc cga ctg tac gtt ggc cgc gta ctg gac aag gat ggc aga ccg				5738
Met Ser Arg Leu Tyr Val Gly Arg Val Leu Asp Lys Asp Gly Arg Pro				
1210		1215	1220	
ctg ctg gac gca cag cca ctg aac tat cca ttt ttg tgg ttg gga cct				5786
Leu Leu Asp Ala Gln Pro Leu Asn Tyr Pro Phe Leu Ser Leu Gly Pro				
1225		1230	1235	
tcc ggg cga ttt agc ctg cag agc gag cat aaa gaa tcc agc ctg tgg				5834
Ser Gly Arg Phe Ser Leu Gln Ser Glu His Lys Glu Ser Ser Leu Trp				
1240		1245	1250	
ctg ctg tct aaa aac agg atc ctg cgt tgt ccg atg tca gta cat aaa				5882
Leu Leu Ser Lys Asn Arg Ile Leu Arg Cys Pro Met Ser Val His Lys				
1255		1260	1265	1270
cgt cgg gat gtt atg cag gta gtg ggt gat gtg cgg tgt gaa tta agt				5930
Arg Arg Asp Val Met Gln Val Val Gly Asp Val Arg Cys Glu Leu Ser				
1275		1280	1285	
gac gtg gat gcc ctg cca cag gcg ttg caa ata tgg ccg cgg gtc atc				5978
Asp Val Asp Ala Leu Pro Gln Ala Leu Gln Ile Ser Pro Arg Val Ile				
1290		1295	1300	
cgt ttg ctg aac gtg gca ggt ttg ctg cgc cat tcc gtt cag gaa gcc				6026
Arg Leu Leu Asn Val Ala Gly Leu Leu Arg His Ser Val Gln Glu Ala				
1305		1310	1315	
tga cgtagagata aaggcgtaa ct atg agt aat aaa atg aag tgg acg agt				6078
Met Ser Asn Lys Met Lys Trp Thr Ser				
1320		1325		
atg aca gcc cat tgg tca gca att att aat ttc atc cga aaa tat gtt				6126
Met Thr Ala His Trp Ser Ala Ile Ile Asn Phe Ile Arg Lys Tyr Val				
1330		1335	1340	
tat cca gca agg ata att gcc atc ctg ctg atg gct ggc gct aca ctg				6174
Tyr Pro Ala Arg Ile Ile Ala Ile Leu Leu Met Ala Gly Ala Thr Leu				
1345		1350	1355	1360
cca caa gtc gcc gat gcg att acc gtc gac ctg aat tac gac aag aac				6222
Pro Gln Val Ala Asp Ala Ile Thr Val Asp Leu Asn Tyr Asp Lys Asn				
1365		1370	1375	
aat gta gcg gtc atc act cct gtc tgg tcc caa gaa tgg agt gta gca				6270
Asn Val Ala Val Ile Thr Pro Val Trp Ser Gln Glu Trp Ser Val Ala				
1380		1385	1390	
aat gtg ttg ggg gga tgg gta tgt cgt tca aac agg aat gaa aat gag				6318
Asn Val Leu Gly Gly Trp Val Cys Arg Ser Asn Arg Asn Glu Asn Glu				
1395		1400	1405	
ggg gcg tgt gaa gaa aca cat ttg gta tgg tgg tat gct ttt gga gct				6366
Gly Ala Cys Glu Glu Thr His Leu Val Trp Trp Tyr Ala Phe Gly Ala				

1410	1415	1420	
tat tca aaa att cgt ctg cgt ttc aga gaa caa atc agc cat gcc gaa			6414
Tyr Ser Lys Ile Arg Leu Arg Phe Arg Glu Gln Ile Ser His Ala Glu			
1425	1430	1435	1440
att acg ctc ata ctg ctc ggc agt gtt cgt gat gcc tgt tat act ggt			6462
Ile Thr Leu Ile Leu Leu Gly Ser Val Arg Asp Ala Cys Tyr Thr Gly			
1445	1450	1455	
gtc atc aac atg aac gct gct gca tgt caa tgg ggt agg tcg ctg aaa			6510
Val Ile Asn Met Asn Ala Ala Ala Cys Gln Trp Gly Arg Ser Leu Lys			
1460	1465	1470	
ctt agg ata cct tca gaa gag ctt gcg aag ata cct aca agc gga aca			6558
Leu Arg Ile Pro Ser Glu Glu Leu Ala Lys Ile Pro Thr Ser Gly Thr			
1475	1480	1485	
tgg aaa gca acg tta gtc ctg gat tat tta caa tgg ggc gga gac gat			6606
Trp Lys Ala Thr Leu Val Leu Asp Tyr Leu Gln Trp Gly Gly Asp Asp			
1490	1495	1500	
cct tta ggc aca tca act aca gat atc acg ctg aat gta aca gac cac			6654
Pro Leu Gly Thr Ser Thr Thr Asp Ile Thr Leu Asn Val Thr Asp His			
1505	1510	1515	1520
ttt gct gaa aat gcg gct att tac ttt ccg caa ttt ggt aca gca acg			6702
Phe Ala Glu Asn Ala Ala Ile Tyr Phe Pro Gln Phe Gly Thr Ala Thr			
1525	1530	1535	
ccc cgg gtg gac ctg aat ctt cac cgg atg aat gcc tca caa atg tcg			6750
Pro Arg Val Asp Leu Asn Leu His Arg Met Asn Ala Ser Gln Met Ser			
1540	1545	1550	
ggc agg gct aat ctg gat atg tgt ctg tat gac gga ggt gtg aaa gcc			6798
Gly Arg Ala Asn Leu Asp Met Cys Leu Tyr Asp Gly Gly Val Lys Ala			
1555	1560	1565	
cgt tca tta cag atg aag ata gaa gga agc aat aag tca ggt acg gga			6846
Arg Ser Leu Gln Met Lys Ile Glu Gly Ser Asn Lys Ser Gly Thr Gly			
1570	1575	1580	
ttt cag gtt ata aag agc gat tct gct gat acg att gat tat gcg gtc			6894
Phe Gln Val Ile Lys Ser Asp Ser Ala Asp Thr Ile Asp Tyr Ala Val			
1585	1590	1595	1600
agt atg aat tat ggg gga cga agt att cct gtc acc cgt ggc gtg gag			6942
Ser Met Asn Tyr Gly Gly Arg Ser Ile Pro Val Thr Arg Gly Val Glu			
1605	1610	1615	
ttc agt ctg gat aac gtg gat aaa gca gca acg cgt ccg gtg gta ctt			6990
Phe Ser Leu Asp Asn Val Asp Lys Ala Ala Thr Arg Pro Val Val Leu			
1620	1625	1630	
ccc ggg caa cgg cag gcg gta cgt tgt gtg cca gtg ccc ctt acc ctg			7038
Pro Gly Gln Arg Gln Ala Val Arg Cys Val Pro Val Pro Leu Thr Leu			
1635	1640	1645	

aca aca caa ccc ttt aac atc aga gag aag cgt tct ggt gag tat cag 7086  
 Thr Thr Gln Pro Phe Asn Ile Arg Glu Lys Arg Ser Gly Glu Tyr Gln  
 1650 1655 1660

gga acg ctg aca gtg aca atg ctg atg gga aca caa acc ccc tga 7131  
 Gly Thr Leu Thr Val Thr Met Leu Met Gly Thr Gln Thr Pro  
 1665 1670 1675

cagtaattat ttattttatt gatattcttc ttatatgggt ttttaaataca gagttctctt 7191  
 tatatacttg ttttatttaa taaagagaat ctattcactt atgaaaatca atgcgtgagg 7251

ttctgctttc ct atg act gtg tat tta gat gat aaa gat aaa gaa tta ttg 7302  
 Met Thr Val Tyr Leu Asp Asp Lys Asp Lys Glu Leu Leu  
 1680 1685 1690

aaa gaa atc caa aaa gat tgt gca caa act tta tgg caa ctt gca tat 7350  
 Lys Glu Ile Gln Lys Asp Cys Ala Gln Thr Leu Trp Gln Leu Ala Tyr  
 1695 1700 1705

aaa gtg gga ctt acg ccc aca cca tgt ttc aaa cgt tta aaa aaa ctt 7398  
 Lys Val Gly Leu Thr Pro Thr Pro Cys Phe Lys Arg Leu Lys Lys Leu  
 1710 1715 1720

aaa gac agg ggg gtt atc att ggt cag ttc gct tta ttg gat aag gaa 7446  
 Lys Asp Arg Gly Val Ile Ile Gly Gln Phe Ala Leu Leu Asp Lys Glu  
 1725 1730 1735 1740

aaa cta ggt ctt tca ctt aat gtc ttt att atg att aac ata tct gag 7494  
 Lys Leu Gly Leu Ser Leu Asn Val Phe Ile Met Ile Asn Ile Ser Glu  
 1745 1750 1755

gag caa tac gct agt att tct gag aaa ata aag tca atg cct gag gtt 7542  
 Glu Gln Tyr Ala Ser Ile Ser Glu Lys Ile Lys Ser Met Pro Glu Val  
 1760 1765 1770

att gcc ttt tat cga att tct gga tca ttt aat tat tta atg cat aca 7590  
 Ile Ala Phe Tyr Arg Ile Ser Gly Ser Phe Asn Tyr Leu Met His Thr  
 1775 1780 1785

gta ttt aca gat atg aac gat tac tat agt ttt tat gag aaa ata ata 7638  
 Val Phe Thr Asp Met Asn Asp Tyr Tyr Ser Phe Tyr Glu Lys Ile Ile  
 1790 1795 1800

tta act aat tct tca att agt gga tct gca tgg agc ttt gtt ctt gag 7686  
 Leu Thr Asn Ser Ser Ile Ser Gly Ser Ala Ser Ser Phe Val Leu Glu  
 1805 1810 1815 1820

caa ata aag gaa aca aac gaa ctg tca gtg tga aagtgtgatg tgtacttact 7739  
 Gln Ile Lys Glu Thr Asn Glu Leu Ser Val  
 1825 1830

gatttaatac attattatcc ttcttacgga acaacaacgg cagattgcgg ctgttgaaca 7799

aggatttttaa tcagcagtggtg tgaaattaag cggcacagaa taacacagcg gaatatcaca 7859

tggttaaata tcaccccggtg catgtaacaa aaaaccgcat taaaacagat gatgttactg 7919

atatttattt cgttgaaccc ttctggaaaa aaggcgaaaa ccacataatt gagtcattga 7979

tggtttttga agagttacaa aagtcattta atttattcaa ccataaatat ggggttaaata 8039  
 aatataact caggatcccc tgggaatttg tgctcataca tatggaaagg atcagtaaata 8099  
 taaatagcgt cgggttattt gctgtttctg ttgactttta taacaaccac aaatttctga 8159  
 gcgagtagat caggagtcgc agagattatg gtatggaagt ttggtttgat ttttgtggta 8219  
 aacattctta ttccagtga attaaaaacc ttggattcct ttttcaggct tgcgtagtgc 8279  
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 tcggggatat aaatgatgta gaacagaggg ccgtgtacca gaacgaagt gattacatgt 8399  
 atggaatgca atggccatcg tcatatgacg gttttttcct tcgggatcat aaaaaaatg 8459  
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 ttgtttaatt attgttttgt tttttattac gattaaatat aaagaacatc attgttcgtg 8579  
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 atacgaatgg tttatatatg cactagatgt attatttttag tttaatatat cgatgggtgc 8699  
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 acactcacat tactaatcat tattaatatg agtggtggtc ttgttttacg catgcatggt 8819  
 tgcattgtgac gttaaattta atgagctga ctgtatgaat tctaaatact ttagagaggt 8879  
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 gatgctactg tagagataat aaattctgct aaagattccc caattcttgt gcattgacat 9059  
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<210> 2  
 <211> 236  
 <212> PRT  
 <213> Salmonella typhi

<400> 2  
 Met Asn Phe Lys Asp Thr Leu Pro Gly Val Phe Leu Cys Val Ala Met  
 1 5 10 15  
 Phe Ala Cys Gly His Ala Arg Ala Asn Met Leu Val Tyr Pro Met Ala  
 20 25 30  
 Ala Glu Ile Asn Ser Ser Arg Glu Glu Ala Thr Ser Leu Phe Val Tyr  
 35 40 45

Ser Lys Ser Asp His Val Gln Tyr Ile Arg Thr Arg Ile Met Arg Ile  
 50 55 60  
 Glu His Pro Gly Met Pro Gln Glu Lys Glu Val Pro Ala Gly Asn Asp  
 65 70 75 80  
 Ile Glu Thr Gly Leu Val Val Ser Pro Glu Lys Phe Ala Leu Ser Pro  
 85 90 95  
 Gly Thr Lys Lys Thr Ile Arg Val Ile Ser Thr Gln Ala Pro Glu Arg  
 100 105 110  
 Glu Glu Ala Trp Arg Val Tyr Phe Glu Ala Val Pro Glu Leu Glu Asp  
 115 120 125  
 Asp Pro Gln Ala Gly Gly Lys Gln Asn Ser Ser Val Ser Val Asn Leu  
 130 135 140  
 Val Trp Gly Val Leu Leu Arg Val Ser Pro Ser Asp Pro Arg Pro Ala  
 145 150 155 160  
 Leu Val Thr Asp Gly His His Leu Leu Asn Thr Gly Asn Thr Arg Leu  
 165 170 175  
 Ser Leu Ile Arg Ala Gly Asn Cys Asp Thr Thr Cys His Trp Gln Asn  
 180 185 190  
 Ile Gly Lys Ser Ile Tyr Pro Gly Gly Ser Ala Asp Ile Pro Ala Gly  
 195 200 205  
 Ile Lys Ser Asn Ala Phe Arg Val Glu Tyr Arg Thr Gly Ala Asn Ser  
 210 215 220  
 Pro Val Ile Ser Ala Asp Leu Thr Ala Ala Gly Lys  
 225 230 235

<210> 3  
 <211> 191  
 <212> PRT  
 <213> Salmonella typhi

<400> 3  
 Met Tyr Thr Glu Cys Thr Tyr Ile Thr Val Ile Asn Asn Lys Ala Arg  
 1 5 10 15  
 Leu Phe Phe Met Asn Met Lys Thr Ser Phe Ile Ala Ala Val Ala  
 20 25 30  
 Leu Ala Thr Val Tyr Ser Phe Ser Val Ser Ala Val Gln Lys Asp Ile  
 35 40 45  
 Thr Val Thr Ala Asn Ile Asp Ser Thr Leu Glu Leu Leu Gln Ala Asp  
 50 55 60  
 Gly Ser Ser Leu Pro Ser Thr Met Lys Leu Asp Phe Met Pro Gly Lys  
 65 70 75 80  
 Gly Leu Val His Lys Ser Leu Gln Thr Arg Leu Tyr Ser Asn Asp Gln  
 85 90 95

Thr Lys Ser Val Asn Val Lys Leu Leu Asn Ala Pro Gln Leu Ile Asn  
 100 105 110  
 Val Leu Asp Pro Thr Lys Thr Ile Asp Met Glu Val Thr Leu Gly Gly  
 115 120 125  
 Arg Ser Leu Thr Thr Thr Asn Ser Val Leu Glu Ala Lys Thr Leu Phe  
 130 135 140  
 Pro Asp Gly Lys Thr Gly Asp Ala Ser Ala Leu Leu Asn Leu Asp Ile  
 145 150 155 160  
 Gly Gln Lys Ala Gly Ala Ala Leu Gln Asn Leu Pro Ala Gly Glu Tyr  
 165 170 175  
 Ser Gly Leu Val Ser Leu Val Ile Ser Gln Ala Val Thr Ala Gly  
 180 185 190

<210> 4  
 <211> 889  
 <212> PRT  
 <213> Salmonella typhi

<400> 4  
 Met Tyr Tyr Leu Leu Gly Leu Cys Ser Phe Thr Ser Gln Ala Thr Leu  
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 Ile Pro Pro Pro Gly Phe Glu Ser Leu Leu Glu Gly Gln Thr Glu Gln  
 20 25 30  
 Ile Glu Val Leu Leu Pro Gly His Ser Leu Gly Leu Phe Pro Val Val  
 35 40 45  
 Val Lys Pro Asp Thr Val Gln Phe Met Ser Pro Leu Met Val Leu Glu  
 50 55 60  
 Ser Ser Gly Leu Ala Ala Leu Pro Ala Ala Glu Arg Gln Lys Ala Leu  
 65 70 75 80  
 Ala Ala Leu Ser Arg Pro Leu Leu Arg Asn Ser Asn Leu Val Cys Gly  
 85 90 95  
 Val Ser Glu Ala Lys Asp Ser Ser Glu Cys Gly Tyr Val Ala Thr Asp  
 100 105 110  
 Lys Glu Asp Val Ala Val Ile Phe Asp Glu Asn Asn Ala Gln Leu Ser  
 115 120 125  
 Leu Phe Leu Asn Arg Asp Trp Leu Pro Asp Glu Glu Arg Arg Asp Lys  
 130 135 140  
 Arg Trp Leu Thr Pro Thr Pro Glu Gly Val Ser Ala Phe Ile His Arg  
 145 150 155 160  
 Gln Thr Leu Tyr Leu Ser Asp Asp Leu His Ser Arg Asn Met Thr Leu  
 165 170 175

Asn Gly Ser Gly Ala Leu Gly Leu Gly Asp Gly Arg Tyr Leu Gly Gly  
180 185 190

Asp Trp Ala Ala Ile Trp Asn Gln Ser Glu His Tyr Asn Asn Ser Gln  
195 200 205

Ala Trp Phe Asp Asn Leu Phe Val Arg Gln Asp Leu Gly Asn Gln Tyr  
210 215 220

Tyr Leu Gln Ala Gly Arg Met Asp Gln Arg Asn Leu Ser Ser Ala Thr  
225 230 235 240

Gly Gly Asp Phe Gly Phe Ser Leu Leu Pro Leu Ser Arg Phe Asp Gly  
245 250 255

Leu Arg Thr Gly Thr Thr Gln Ala Tyr Val Asn His Glu Val Asp His  
260 265 270

Asn Ala Thr Pro Val Met Val Gln Val Thr Arg Asn Ala Arg Ile Asp  
275 280 285

Ile Tyr Arg Gly Ser Glu Leu Leu Gly Ser Gln Phe Leu Thr Pro Gly  
290 295 300

Met His Thr Leu Asp Thr His Ser Leu Pro Pro Gly Ser Tyr Pro Leu  
305 310 315 320

Ala Leu Arg Val Tyr Glu Asp Gly Ile Leu Arg Arg Thr Glu Thr Gln  
325 330 335

Pro Phe Ser Lys Gly Gly Asn Ser Phe Ser Ala Gln Thr Gln Trp Phe  
340 345 350

Ile Gln Gly Gly Leu Glu Asp Thr Gly Asp Lys Ala Ser His Tyr Asp  
355 360 365

Gly Glu Thr Val Met Ala Ala Gly Phe Gln Thr Gly Leu Arg Lys Asn  
370 375 380

Ile Ser Leu Thr Glu Gly Ile Ser Leu Ala His Glu Ala Trp Tyr Ser  
385 390 395 400

Glu Thr Arg Leu Asn Ser Gln His Ala Val Leu Asp Gly Thr Leu Asp  
405 410 415

Leu Ser Ala Gly Ile Leu His Gly Thr Asp Ser Thr Ser Gly Asn Thr  
420 425 430

Glu Gln Val Thr Tyr Asn Asp Gly Phe Ser Ala Ser Leu Trp Arg Asn  
435 440 445

His Thr Glu Ser Asp Ala Cys Ser Gly Arg His Pro Gln Ser Val His  
450 455 460

Ala Ser Met Thr Cys Gln Thr Ser Met Asn Ala Ser Leu Ser Val Ser  
465 470 475 480

Val Gly Asn Trp Tyr Ala Leu Leu Gly Tyr Ser Thr Ser Arg Thr Glu  
485 490 495

Gly Arg Pro Val Tyr Arg Gly Tyr Asp Asp Asn Ser Asp Lys Glu Asn  
 500 505 510  
 Val Phe Trp Arg Gln Ala Tyr Ile Pro Ala Ser His Arg Glu Ser Ala  
 515 520 525  
 Gln Ala Ser Ala Thr Tyr Ser Leu Asn Met Ala Gly Met Asn Ile Asn  
 530 535 540  
 Thr His Gly Gly Val Trp Arg Thr Arg Asn Asp Gly Val Asn Asp Asp  
 545 550 555 560  
 Gly Leu Phe Met Ser Val Ser Val Ser Tyr Ala Ser Gln Pro Pro Thr  
 565 570 575  
 Met Thr Gly Ser Asn Arg Tyr Thr Ser Ala Gly Thr Asp Ile His Ser  
 580 585 590  
 Ser Arg Asn Gln Lys Thr Gln Thr Ser Trp Asn Val Asn His Val Arg  
 595 600 605  
 Ser Trp Gln Gln Asp Leu Tyr Arg Glu Leu Ser Val Gly Phe Ser Gly  
 610 615 620  
 Tyr Asn Asp Asp Ser Trp Ser Gly Ser Leu Gly Gly Arg Met Ser Gly  
 625 630 635 640  
 Arg Met Gly Glu Leu Ser Ala Thr Ile Ser Asn Ser His Gln Arg Asn  
 645 650 655  
 Ala Gly Ser Ala Ser Ser Leu Thr Ala Gly Tyr Ser Ser Ser Leu Ala  
 660 665 670  
 Leu Ser Arg Asn Gly Leu Phe Trp Gly Gly Gly Gln Asp Gly Glu Pro  
 675 680 685  
 Ala Ser Gly Met Ala Val Asn Val Glu Ser Glu Gly Asp Glu Gly Ser  
 690 695 700  
 Ser Gly Lys Val Val Ser Val Arg Gly Ser Ser Gln Pro Phe Ser Leu  
 705 710 715 720  
 Gly Phe Gly Gln Gln Ser Leu Leu Leu Met Glu Gly Tyr Asn Ala Thr  
 725 730 735  
 Glu Val Thr Ile Glu Asp Ala Gly Val Ser Ser Gln Gly Met Ala Gly  
 740 745 750  
 Val Lys Ala Gly Gly Gly Ser Arg Cys Tyr Phe Leu Thr Pro Gly His  
 755 760 765  
 Leu Leu Val His Asn Ile Ser Ala Ser Met Ser Arg Leu Tyr Val Gly  
 770 775 780  
 Arg Val Leu Asp Lys Asp Gly Arg Pro Leu Leu Asp Ala Gln Pro Leu  
 785 790 795 800  
 Asn Tyr Pro Phe Leu Ser Leu Gly Pro Ser Gly Arg Phe Ser Leu Gln  
 805 810 815



Ser Glu His Lys Glu Ser Ser Leu Trp Leu Leu Ser Lys Asn Arg Ile  
820 825 830  
Leu Arg Cys Pro Met Ser Val His Lys Arg Arg Asp Val Met Gln Val  
835 840 845  
Val Gly Asp Val Arg Cys Glu Leu Ser Asp Val Asp Ala Leu Pro Gln  
850 855 860  
Ala Leu Gln Ile Ser Pro Arg Val Ile Arg Leu Leu Asn Val Ala Gly  
865 870 875 880  
Leu Leu Arg His Ser Val Gln Glu Ala  
885

<210> 5  
<211> 359  
<212> PRT  
<213> Salmonella typhi

<400> 5  
Met Ser Asn Lys Met Lys Trp Thr Ser Met Thr Ala His Trp Ser Ala  
1 5 10 15  
Ile Ile Asn Phe Ile Arg Lys Tyr Val Tyr Pro Ala Arg Ile Ile Ala  
20 25 30  
Ile Leu Leu Met Ala Gly Ala Thr Leu Pro Gln Val Ala Asp Ala Ile  
35 40 45  
Thr Val Asp Leu Asn Tyr Asp Lys Asn Asn Val Ala Val Ile Thr Pro  
50 55 60  
Val Trp Ser Gln Glu Trp Ser Val Ala Asn Val Leu Gly Gly Trp Val  
65 70 75 80  
Cys Arg Ser Asn Arg Asn Glu Asn Glu Gly Ala Cys Glu Glu Thr His  
85 90 95  
Leu Val Trp Trp Tyr Ala Phe Gly Ala Tyr Ser Lys Ile Arg Leu Arg  
100 105 110  
Phe Arg Glu Gln Ile Ser His Ala Glu Ile Thr Leu Ile Leu Leu Gly  
115 120 125  
Ser Val Arg Asp Ala Cys Tyr Thr Gly Val Ile Asn Met Asn Ala Ala  
130 135 140  
Ala Cys Gln Trp Gly Arg Ser Leu Lys Leu Arg Ile Pro Ser Glu Glu  
145 150 155 160  
Leu Ala Lys Ile Pro Thr Ser Gly Thr Trp Lys Ala Thr Leu Val Leu  
165 170 175  
Asp Tyr Leu Gln Trp Gly Gly Asp Asp Pro Leu Gly Thr Ser Thr Thr  
180 185 190  
Asp Ile Thr Leu Asn Val Thr Asp His Phe Ala Glu Asn Ala Ala Ile  
195 200 205

Tyr Phe Pro Gln Phe Gly Thr Ala Thr Pro Arg Val Asp Leu Asn Leu  
 210 215 220  
 His Arg Met Asn Ala Ser Gln Met Ser Gly Arg Ala Asn Leu Asp Met  
 225 230 235 240  
 Cys Leu Tyr Asp Gly Gly Val Lys Ala Arg Ser Leu Gln Met Lys Ile  
 245 250 255  
 Glu Gly Ser Asn Lys Ser Gly Thr Gly Phe Gln Val Ile Lys Ser Asp  
 260 265 270  
 Ser Ala Asp Thr Ile Asp Tyr Ala Val Ser Met Asn Tyr Gly Gly Arg  
 275 280 285  
 Ser Ile Pro Val Thr Arg Gly Val Glu Phe Ser Leu Asp Asn Val Asp  
 290 295 300  
 Lys Ala Ala Thr Arg Pro Val Val Leu Pro Gly Gln Arg Gln Ala Val  
 305 310 315 320  
 Arg Cys Val Pro Val Pro Leu Thr Leu Thr Thr Gln Pro Phe Asn Ile  
 325 330 335  
 Arg Glu Lys Arg Ser Gly Glu Tyr Gln Gly Thr Leu Thr Val Thr Met  
 340 345 350  
 Leu Met Gly Thr Gln Thr Pro  
 355

<210> 6  
 <211> 151  
 <212> PRT  
 <213> Salmonella typhi

<400> 6  
 Met Thr Val Tyr Leu Asp Asp Lys Asp Lys Glu Leu Leu Lys Glu Ile  
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 Gln Lys Asp Cys Ala Gln Thr Leu Trp Gln Leu Ala Tyr Lys Val Gly  
 20 25 30  
 Leu Thr Pro Thr Pro Cys Phe Lys Arg Leu Lys Lys Leu Lys Asp Arg  
 35 40 45  
 Gly Val Ile Ile Gly Gln Phe Ala Leu Leu Asp Lys Glu Lys Leu Gly  
 50 55 60  
 Leu Ser Leu Asn Val Phe Ile Met Ile Asn Ile Ser Glu Glu Gln Tyr  
 65 70 75 80  
 Ala Ser Ile Ser Glu Lys Ile Lys Ser Met Pro Glu Val Ile Ala Phe  
 85 90 95  
 Tyr Arg Ile Ser Gly Ser Phe Asn Tyr Leu Met His Thr Val Phe Thr  
 100 105 110

Asp Met Asn Asp Tyr Tyr Ser Phe Tyr Glu Lys Ile Ile Leu Thr Asn  
115 120 125

Ser Ser Ile Ser Gly Ser Ala Ser Ser Phe Val Leu Glu Gln Ile Lys  
130 135 140

Glu Thr Asn Glu Leu Ser Val  
145 150

1  
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Claims:

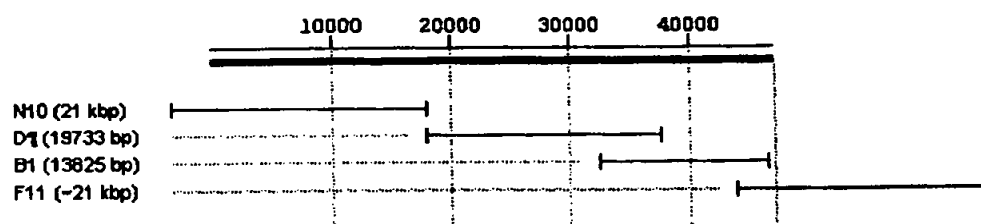
1. Protein encoded by a nucleotide sequence selected from the group consisting of SEQ ID NO 1 and SEQ ID NO 2, or parts thereof, for use in medicine.
2. Antibodies directed against the protein encoded by a nucleotide sequence selected from the group consisting of SEQ ID NO 1 and SEQ ID NO 2, or antigenic fragments thereof for use in medicine.
3. Nucleotide sequence selected from the group consisting of SEQ ID NO 1 and SEQ ID NO 2, or parts thereof, for use in medicine.
4. A vaccine for the protection against diseases caused by *Salmonella enterica* subspecies I, comprising the protein, or parts thereof, encoded by the nucleotide sequence according to SEQ ID NO 1 or antibodies directed against the protein encoded by SEQ ID NO 1, or antigenic fragments thereof and, optionally, a pharmaceutically acceptable carrier.
5. A vaccine for the protection against diseases caused by *Salmonella enterica* subspecies I serovar Typhi, comprising the protein, or parts thereof, encoded by the nucleotide sequence according to SEQ ID NO 2 or antibodies directed against the protein encoded by SEQ ID NO 2, or antigenic fragments thereof and, optionally, a pharmaceutically acceptable carrier.
6. A nucleic acid vaccine for the protection against diseases caused by *Salmonella enterica* subspecies I, comprising SEQ ID NO 1, or parts thereof and, optionally, a pharmaceutically acceptable carrier.
7. A nucleic acid vaccine for the protection against diseases caused by *Salmonella enterica* subspecies I serovar Typhi, comprising SEQ ID NO 2, or parts thereof and, optionally, a pharmaceutically acceptable carrier.
8. A vector vaccine for the protection against diseases caused by *Salmonella enterica* subspecies I, comprising a host in which a recombinant vector comprising SEQ ID NO 1, or parts thereof, has been inserted and, optionally, a pharmaceutically acceptable carrier.

9. A vector vaccine for the protection against diseases caused by *Salmonella enterica* subspecies I serovar Typhi, comprising a host in which a recombinant vector comprising SEQ ID NO 2, or parts thereof, has been inserted and,  
5 optionally, a pharmaceutically acceptable carrier.
10. A method for protection against diseases caused by *Salmonella enterica* subspecies I, comprising administering a vaccine according to any of claims 4, 6, and 8.  
10
11. A method for protection against diseases caused by *Salmonella enterica* subspecies I serovar Typhi, comprising administering a vaccine according to any of claims 5, 7, and 9.
12. Antibodies directed against the protein encoded by a nucleotide sequence selected from the group consisting of SEQ ID NO 1 and SEQ ID NO 2, or antigenic fragments thereof, for use in a diagnostic method.  
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13. Protein encoded by a nucleotide sequence selected from the group consisting of SEQ ID NO 1 and SEQ ID NO 2, or parts thereof, for use in a diagnostic method.  
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14. Primers for or, probes that hybridize with a nucleotide sequence selected from the group consisting of SEQ ID NO 1 and SEQ ID NO 2, for use in a  
25 diagnostic method.

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## ABSTRACT

5 The present invention is based on the finding that two fimbrial structures are specific for *Salmonella enterica* subspecies 1 bacteria. Due to their specificity they can be used to provide vaccines against *Salmonella enterica* subspecies I as well as for detection of *Salmonella enterica* subspecies I.



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